AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Original) Compounds with general formula (I)

in which

m is the number 0 or 1;

Z and Z', which can be the same or different, are an integer ranging from 0 to 2;

Y and Y', which can be the same or different, are $(CH_2)n_1$; $(CH_2)n_2$ - $CH[NR^{VII}(CH_2)n_4-NHR^I]$ - $(CH_2)n_3$; CH_2 - $CH[CH_2$ - $CH_2]_2$ - or $(CH_2)n_2$ - $N[(CH_2)n_4$ - $NHR^{IV}]$ - $(CH_2)n_3$;

GIANNINI et al.

New*National Phase Application Based on PCT/IT2004/000374

Y" is selected from the group consisting of H; cycloalkyl C_3 - $C_{7;}$ (CH₂) n_5 -N[CH₂-CH₂]₂N-(CH₂) n_6 NHR^V; (CH₂) n_7 -CH[CH₂-CH₂]₂NR^V;

X is O, or is a simple bond;

n-n₈, which can be the same or different, are an integer ranging from 0 to 5;

 R^{I} , R^{III} , R^{IV} , and R^{V} , which can be the same or different, are a protective group for the nitrogen to which they are bound; CO_2R^{VI} ; CO_2CH_2Ar ; $CO_2(9-fluorenylmethyl)$; $(CH_2)n_5-NHCO_2R^{VI}$; CH_2Ar ; COAr; $(CH_2)n_5-NHCO_2CH_2Ar$; $(CH_2)n_5-NHCO_2-(9-fluorenylmethyl)$.

R^{VI} is a straight or branched (C₁-C₆) alkyl;

R^{VII} is H or R^I-R^V;

Ar is a C_6 - C_{12} aromatic residue, such as phenyl, optionally substituted with one or more groups selected from: halogen, hydroxy, C_1 - C_5 alkyl, C_1 - C_5 alkoxy, phenyl, cyano, nitro, -NR^{VIII}R^{IX}, where R^{VIII} and R^{IX}, which can be the same or different, are hydrogen, straight or branched (C_1 - C_5) alkyl, or Ar is a heterocyclic group, said heterocyclic group containing at least one heteroatom selected from a nitrogen atom, optionally substituted with a (C_1 - C_5) alkyl group, and/or oxygen and/or sulphur; said heterocycle can be substituted with one or more groups selected from halogen, hydroxy, C_1 - C_5 alkyl, C_1 - C_5 alkoxy, phenyl, cyano, nitro, -NR^{VIII}R^{IX}, where R^{VIII} and R^{IX}, which can be the same or different, are hydrogen, straight or branched (C_1 - C_5) alkyl, the N₁-oxides, racemic mixtures, their individual enantiomers, their individual diastereoisomers, the *E* and *Z* forms, their mixtures, and pharmaceutically acceptable salts.

GIANNINI et al.

New National Phase Application Based on PCT/IT2004/000374

- 2. (Original) Compounds according to claim 1, in which the protective groups are bulky groups of a lipophilic nature.
- 3. (Original) Compounds according to claim 1, in which the protective groups are selected from the group consisting of: CO_2R^{VI} ; CO_2CH_2Ar ; CO_2 -(9-fluorenylmethyl); $(CH_2)n_5$ -NHCO₂R^{VI}; $(CH_2)n_5$ -NHCO₂CH₂Ar; $(CH_2)n_5$ -NHCO₂-(9-fluorenylmethyl), in which R^{VI} is as defined above.
- 4. (Original) Compounds according to claim 3, in which the protective groups are selected from the group consisting of tert-butoxycarbonyl; benzyloxycarbonyl; 9-fluorenylmethyloxycarbonyl.
- 5. (Currently Amended) Compounds according to any of claimsclaim 1[[-4]], in which m is 0.
- 6. (Original) Compounds according to claim 5, selected from the group consisting of:
- tert-butylester of 20S-(4-{[3-(7-camptothecinylidene-amino)-propyl]-tert-butoxycarbonyl-amino}-butyl)-(3-tert-butoxycarbonylaminopropyl)-carbamic acid;
- tert-butylester of 20S-(4-{[3-(7-camptothecinylidene-amino)-propyl]-tert-butoxycarbonyl-amino}-butyl)-carbamic acid;
- tert-butylester of 20S-[3-(7-camptothecinylidene-amino)-butyl]-carbamic acid;
- 20S-7-[3-(N-tert-butoxycarbonylamino)propoxyimino-methyl]-camptothecin.

GIANNINI et al. New National Phase Application Based on PCT/IT2004/000374

- 7. (Currently Amended) Compounds according to any of claimsclaim 1[[-4]], in which m is 1.
- 8. (Original) Compounds according to claim 7, selected from the group consisting of:
- tert-butylester of 20RS-(4-{[3-(7-homocamptothecinylidene-amino)-propyl]-tert-butoxycarbonyl-amino}-butyl)-(3-tert-butoxyicarbonylaminopropyl)-carbamic acid;
- tert-butylester of 20RS-(4-{[3-(7-homocampto-thecinylidene-amino)-propyl]-tert-butoxycarbonyl-amino}-butyl)-carbamic acid;
- tert-butylester of 20RS-[3-(7-homocamptothecinylidene-amino)-butyl]-carbamic acid;
- 20R,S-7-[3-(N-tert-butoxycarbonylamino)propoxyimino-methyl]-homocamptothecin
- 9. (Currently Amended) Pharmaceutical composition containing at least one compound according to claims claim 1-8 as the active ingredient in admixture with at least one pharmaceutically acceptable vehicle and/or excipient.
- 10. (Currently Amended) Use of compounds according to elaims claim 1-8 as medicaments.

GIANNINI et al.

New National Phase Application Based on PCT/IT2004/000374

- 11. (Currently Amended) Use of compounds according to <u>claims_claim_1-8</u> for the preparation of a medicament with topoisomerase 1 inhibiting activity.
- 12. (Original) Use according to claim 11 for the preparation of a medicament with anticancer activity.
- 13. (Original) Use according to claim 11 for the preparation of a medicament with antiparasite activity.
- 14. (Original) Use according to claim 11 for the preparation of a medicament with antiviral activity.